## Amendments to the Claims

- 1. (Cancelled)
- 2. (Currently amended) A compound of formula 1

$$R_{4}$$
  $R_{4}$   $R_{4$ 

wherein  $R_1$ - $R_2$  is a radical of formula  $\boldsymbol{2}$ 

$$R_{5} \longrightarrow N \longrightarrow N \longrightarrow R_{6}$$

wherein R<sub>2</sub> is hydrogen, alkyl of 1 to 10 carbon atoms, or a saccharide moiety; R<sub>5</sub> is hydrogen, halogen, trifluoromethyl, or hydroxy; and R<sub>6</sub> is hydrogen, hydroxy or unsubstituted or substituted amino; and tautomeric forms thereof;

X is oxygen or sulfur;

R<sub>3</sub> is triazolylene, tetrazolylene, isoxazolylene, thienylene, isoxazolidinylene, or alkynylene, wherein a double bond or the triple bond, respectively, is connected to CH<sub>2</sub>;

R<sub>4</sub> is an optionally substituted straight or branched chain alkylene group with 1 to 300 carbon atoms, wherein optionally

- (a) one or more carbon atoms are replaced by oxygen
- (b) one or more carbon atoms are replaced by nitrogen carrying a hydrogen atom, and an adjacent carbon atom is substituted by oxo, representing an amide function –NH-CO-;
- (c) one or more carbon atoms are replaced by oxygen, and an adjacent carbon atom is substituted by oxo, representing an ester function –O-CO-;

- (d) the bond between two adjacent carbon atoms is a double or a triple bond, representing a function -CH=CH- or -CEC-;
- (e) one or more carbon atoms are replaced by a phenylene, a saturated or unsaturated cycloalkylene, a saturated or unsaturated bicycloalkylene, a bridging heteroaromatic or a bridging saturated or unsaturated heterocyclyl group; and/or
- (f) two adjacent carbon atoms are replaced by a disulfide linkage -S-S-; and L is one or a plurality of same or different labels selected from a spectroscopic probe selected from a fluorophore and a chromophore, a moiety which is comprising one part of a specific binding pair selected from biotin, avidin, avidin and streptavidin, an amine, an activated carboxy group, an azide and a propiolic acid derivative, a moiety which is capable of generating hydroxyl radicals upon exposure to H<sub>2</sub>O<sub>2</sub> and ascorbate, a moiety which is capable of generating reactive radicals upon irradiation with light, a moiety covalently attached to a solid support, a nucleic acid moiety or a derivative thereof capable of undergoing base-pairing with its complementary strand, a lipid or other hydrophobic moiety with membrane-inserting properties, a bond connecting R<sub>4</sub> to R<sub>1</sub> forming a cyclic substrate, and a further group -R<sub>3</sub>-CH<sub>2</sub>-X-R<sub>1</sub>-R<sub>2</sub>

## 3. (Cancelled)

- 4. (Previously presented) The compound of formula 1 according to claim 2, wherein the saccharide moiety  $R_2$  is a  $\beta$ -D-2'-deoxyribosyl, or a  $\beta$ -D-2'-deoxyribosyl being incorporated into a single stranded oligodeoxyribonucleotide having a length of 2 to 99 nucleotides, wherein the radical of formula 2 occupies any position within the oligonucleotide sequence.
- 5. (Previously presented) The compound of formula 1 according to claim 2, wherein  $R_2$  is hydrogen,  $R_5$  is hydrogen,  $R_6$  is unsubstituted amino, and X is oxygen.
- 6. (Withdrawn) The compound of formula 1 according to claim 1, wherein  $R_1$ - $R_2$  is a radical of formula 3

$$R_2$$
  $R_6$   $R_6$ 

wherein  $R_2$  is hydrogen, alkyl of 1 to 10 carbon atoms, or a saccharide moiety; and  $R_6$  is hydrogen, hydroxy or unsubstituted or substituted amino; and tautomeric forms thereof.

7. (Withdrawn) The compound of formula 1 according to claim 1, wherein  $R_1$ - $R_2$  is a radical of formula 4

wherein  $R_2$  is hydrogen, alkyl of 1 to 10 carbon atoms, or a saccharide moiety; and  $R_7$  and  $R_8$  are both independently of one another hydrogen, halogen, lower alkyl with 1 to 4 carbon atoms, amino, or nitro.

- 8. (Previously presented) The compound of formula 1 according to claim 2, wherein R<sub>3</sub> is triazolylene, tetrazolylene, isoxazolylene, thienylene, or isoxazolidinylene.
- 9. (Previously presented) The compound of formula 1 according to claim 8 wherein  $R_3$  is triazolylene.
- 10. (Previously presented) The compound of formula 1 according to claim 8 wherein  $R_3$  is tetrazolylene.

- 11. (Previously presented) The compound of formula 1 according to claim 8 wherein R<sub>3</sub> is isoxazolylene.
- 12. (Previously presented) The compound of formula 1 according to claim 8 wherein  $R_3$  is thienylene.
- 13. (Previously presented) The compound of formula 1 according to claim 8 wherein R<sub>3</sub> is isoxazolidinylene.
- 14. (Previously presented) The compound of formula 1 according to claim 2, wherein  $R_3$  is 1-alkynylene.
- 15. (Previously presented) The compound of formula 1 according to claim 2, wherein  $R_4$  is a straight chain alkylene group with 2 to 25 carbon atoms, a straight chain polyethylene glycol group with 4 to 100 ethyleneoxy units, or a straight chain alkylene group with 2 to 25 carbon atoms wherein two or more carbon atoms are replaced by an amide function –NH-CO-, optionally attached to the group  $R_3$  by a –CH=CH- or –CEC- group.
- 16. (Previously presented) The compound of formula 1 according to claim 2, wherein R<sub>4</sub> is a branched chain alkylene group comprising a polyethylene glycol group of 3 to 6 ethylene glycol units and one or more alkylene groups wherein carbon atoms are replaced by amide bonds, and further carrying substituted amino and hydroxy functions.
- 17. (Previously presented) The compound of formula 1 according to claim 2, wherein R<sub>4</sub> is a branched chain alkylene group, wherein amine, carboxamide and ether functions replace carbon atoms of the alkylene group.
- 18. (Previously presented) The compound of formula 1 according to claim 2, wherein L is a further group  $-R_3-CH_2-X-R_1-R_2$ .

- 19. (Previously presented) The compound of formula 1 according to claim 2, wherein R<sub>4</sub> is a straight chain alkylene group of 10 to 40 carbon atoms wherein 3 to 12 carbon atoms are replaced by oxygen, one or two carbon atoms are replaced by 1,4-triazolidene units, and optionally one carbon atom is replaced by a 1,4-phenylene unit.
- 20. (Previously presented) The compound of formula 1 according to claim 2, wherein  $R_4$  is a straight chain alkylene group of 10 to 40 carbon atoms optionally substituted by oxo wherein 3 to 12 carbon atoms are replaced by oxygen and one or two carbon atoms are replaced by nitrogen.
- 21. (Previously presented) The compound of formula 1 according to claim 2, wherein R<sub>4</sub> is a straight chain alkylene group of 6 to 40 carbon atoms wherein 2 to 12 carbon atoms are replaced by oxygen and one or two bonds between two adjacent carbon atoms is a double bond.
- 22. (Previously presented) The compound of formula 1 according to claim 2, wherein  $R_6$  is amino and L is a bond connecting  $R_4$  to  $R_6$ .
- 23. (Previously presented) The compound of formula 1 according to claim 2, wherein L is methotrexate.
- 24. (Previously presented) The compound of formula 1 according to claim 2, wherein L is a plurality of same or different labels.
- 25. (Previously presented) The compound of formula 1 according to claim 24, wherein L is two different labels.
- 26. (Cancelled)
- 27. (Previously presented) A method for the synthesis of a compound of the formula 1 according to claim 2, which comprises reacting a compound of the formula  $R_2$ - $R_1$ -X- $CH_2$ - $R_3$ - $R_4$ ', wherein  $R_1$ ,  $R_2$ ,  $R_3$  and X have the meaning as defined in claim 2 and  $R_4$ ' is a polyfunctional

residue having two or more reactive nucleophilic or electrophilic groups, with a suitable reagent introducing one or more labels L.

- 28. (Currently amended) A method according to claim 27 wherein the reactive nucleophilic or electrophilic groups in  $R_4$  are protected by separately removable protection groups, the method comprising the further steps of
- (a) separately deprotecting one protected reactive nucleophilic or electrophilic group and attaching a label to it or extending the linker polyfunctional residue R<sub>4</sub>,
- (b) separately deprotecting another protected reactive nucleophilic or electrophilic group and attaching a label to it or extending the <u>linker\_polyfunctional residue</u>  $R_4$ , and
- (c) <u>optionally</u> repeating the steps of deprotection and label attachment or <u>linker\_polyfunctional</u> residue extension <u>until all\_depending on the number of protected reactive nucleophilic and electrophilic groups are removed.</u>
- 29. (Previously presented) A compound of the formula 1

$$CH_2-R_3$$
 $X$ 
 $R_1$ 
 $R_2$ 
 $R_1$ 

wherein R<sub>1</sub>-R<sub>2</sub> is a radical of formula 2

$$R_5$$
 $N$ 
 $N$ 
 $R_6$ 
 $R_2$ 

wherein R<sub>2</sub> is hydrogen, R<sub>5</sub> is hydrogen and R<sub>6</sub> is unsubstituted amino;

X is oxygen;

R<sub>3</sub> is triazolylene, tetrazolylene, isoxazolylene, thienylene, isoxazolidinylene or alkynylene, wherein a double bond or the triple bond, respectively, is connected to CH<sub>2</sub>;

R<sub>4</sub> is an optionally substituted straight or branched chain alkylene group with 1 to 300 carbon atoms, wherein optionally

- (a) one or more carbon atoms are replaced by oxygen
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- (c) one or more carbon atoms are replaced by oxygen, and an adjacent carbon atom is substituted by oxo, representing an ester function –O-CO-;
- (d) the bond between two adjacent carbon atoms is a double or a triple bond, representing a function -CH=CH- or -CEC-;
- (e) one or more carbon atoms are replaced by a phenylene, a saturated or unsaturated cycloalkylene, a saturated or unsaturated bicycloalkylene, a bridging heteroaromatic or a bridging saturated or unsaturated heterocyclyl group; and/or
- (f) two adjacent carbon atoms are replaced by a disulfide linkage -S-S-; and L is amino or azido.
- 30. (Previously presented) A compound according to claim 29 wherein R<sub>4</sub> is a straight chain alkylene group of 10 to 40 carbon atoms optionally substituted by oxo wherein up to 12 carbon atoms are replaced by oxygen and zero, one or two carbon atoms are replaced by nitrogen.
- 31. (Previously presented) A method for detecting a protein of interest, which comprises contacting an AGT fusion protein comprising the protein of interest with an AGT substrate carrying a label, and detecting the AGT fusion protein using the label in a system designed for recognising the label, and wherein the AGT substrate carrying the label is a compound of formula 1 according to claim 2.